AMENDMENTS

IN THE CLAIMS:

Please amend claims as set forth below.

Claims 1-38. (Canceled).

- 39. (Currently amended) A method for constructing a synthetic polynucleotide from which a polypeptide is producible to confer a selected phenotype upon an organism of interest or part-thereof an immune response to a target antigen in a mammal of interest in a different quality than that conferred by a parent polynucleotide that encodes the same polypeptide, the method comprising: (a) selecting a first codon of the parent polynucleotide for replacement with a synonymous codon, wherein the synonymous codon is selected on the basis that it exhibits a different phenotypie preference for conferring an immune response than the first codon in a comparison of phenotypie immune response preferences in test organisms mammals or parts thereof, wherein the test organisms mammals are selected from the group consisting of organisms mammals of the same species as the organism mammal of interest and organisms mammals of a different species than the mammal that are related to the organism of interest; and (b) replacing the first codon with the synonymous codon to construct the synthetic polynucleotide, wherein the selected phenotype is other than a phenotype conferred upon a cell by a polynucleotide that is expressed in the cell and that encodes the polypeptide.
- 40. (Currently amended) A method according to claim 39, wherein the phenotypie immune response preferences of codons in the test organisms mammals or parts are compared by: (i) separately introducing into the test organisms mammals or parts individual synthetic constructs, each of which comprises a regulatory polynucleotide operably linked to a tandem repeat of a codon fused in frame with a reporter polynucleotide that encodes a reporter protein, which produces, or which is predicted to produce, a corresponding phenotype an immune response selected from the group consisting of the selected phenotype immune response to the target antigen and an phenotype an immune response of the same class as the selected phenotype the immune response to the target antigen; and (ii) comparing the quality of the phenotype the immune response to the target antigen; and (ii) comparing the quality of the phenotype the immune response immune response preferences of the codons.
- (Currently amended) A method according to claim 39 40, wherein the reporter protein produces the selected-phenotype immune response to the target antigen.

- 42. (Currently amended) A method according to claim 39 <u>40</u>, wherein the reporter protein does not produce the selected phenotype <u>immune response to the target antigen</u> but produces the same class of phenotype as the selected phenotype an <u>immune response of the same class</u> as the <u>immune response to the target antigen</u>.
- 43. (Previously presented) A method according to claim 41 or claim 42, wherein the reporter protein is selected from antigens derived from pathogenic organisms, cancer antigens, self antigens, transplantation antigens, growth factors, hormones and toxins.
- 44. (Currently amended) A method according to claim 39, wherein the phenotype is selected from immunity, <u>and</u> antigen tolerance, angiogenesis, anti-angiogenesis, amelioration of elinical symptoms, reduced or increased cell death, reduced or increased cell differentiation, reduced or increased cell-proliferation, tumour or cancer regression, growth and repair of tissue or organ, decreased fibrosis, inhibition or reversal of cell-senescence, increased or reduced cell migration, differential expression of protein between different cells or tissues of an organism or part thereof, trauma recovery, recovery from burns, antibiotic resistance or sensitivity, herbicide tolerance or sensitivity, starch biosynthesis or modification, fatty acid biosynthesis, disease resistance or tolerance, pest resistance or tolerance including insect resistance or tolerance, viral resistance or tolerance, fungal resistance or tolerance, a metabolic trait including sucross metabolism, frost resistance or tolerance, stress tolerance, and improved food content or increased vields.
 - 45. (Canceled).
- 46. (Currently amended) A method according to claim 45 39, wherein the immune response is a humoral immune response.
- 47. (Currently amended) A method according to claim 45 39, wherein the immune response is a cell mediated immune response.
- 48. (Currently amended) A method according to claim 45 39, wherein the immune response is an innate immunity mediated response.
- 49. (Currently amended) A method according to claim 39 40, wherein the synthetic constructs are introduced into the test organisms mammals using the same or similar mode of introduction.

- 50. (Currently amended) A method according to claim 39 40, wherein the synthetic constructs are introduced into the test organisms mammals at the same or corresponding site.
- 51. (Currently amended) A method according to claim 39 40, wherein the organism of interest is a mammal and the synthetic constructs are introduced by oral, intravenous, intramuscular, intransaal, buccal, subcutaneous, transdermal, buccal or sublingual route.
- 52. (Currently amended) A method according to claim 39 40, wherein the synthetic constructs are introduced into one or more of-eell or tissue types of the organism of interest test mammals.
- 53. (Currently amended) A method according to claim 52, wherein the synthetic constructs are introduced into cells selected from muscle cells, and skin cells, brain cells, lung cells, kidney cells, pancreas cells, cells of a reproductive organ, heart-cells, vascular cells, liver cells, flower cells, meristematic cells, root-cells and leaf cells.
- 54. (Currently amended) A method according to claim 39 40, wherein the tandem repeat of each of the synthetic constructs comprises at least three copies of the corresponding codon.
- 55. (Currently amended) A method according to claim 39, wherein the synonymous codon is selected such that it has a higher phenotypie <u>immune response</u> preference than the first codon.
- 56. (Currently amended) A method according to claim 17 39, wherein the synonymous codon is selected when the quality of the phenotype immune response conferred by the synthetic construct comprising a tandem repeat of the synonymous codon is at least about 5% higher than the quality of the phenotype immune response conferred by the synthetic construct comprising a tandem repeat of the first codon.
- 57. (Currently amended) A method according to claim 39, wherein the synonymous codon is selected such that it has a lower phenotypie immune response preference than the first codon.
- 58. (Currently amended) A method according to claim 57, wherein the synonymous codon is selected when the quality-of-the-phenotype immune response conferred by the synthetic construct comprising a tandem repeat of the synonymous codon is at least about 5% lower than the quality-of-the-phenotype immune response conferred by the synthetic construct comprising a tandem repeat of the first codon.

59. - 65 (Canceled).

66. (Withdrawn-Currently amended) A method for determining the phenotypie immune response preference of a first codon in an organism a mammal of interest or part thereof, the method comprising: (a) introducing a synthetic construct into a test organism mammal or part thereof, wherein the test organism mammal is selected from the group consisting of an organism a mammal of the same species as the organism mammal of interest and an organism a mammal of a different species than that is related to the organism mammal of interest, the synthetic construct comprising a regulatory polynucleotide operably linked to a tandem repeat of the first codon fused in frame with a reporter polynucleotide that encodes a reporter protein, which produces, or which is predicted to produce, an immune response to a target antigen selected phenotype or an immune response phenotype of the same class as the immune response to a target antigen selected phenotype; and (b) determining the quality of the corresponding phenotype immune response displayed by the test mammal organism or part, wherein the selected phenotype or the phenotype of the same class as the selected phenotype is other than a phenotype conferred upon a cell by a polynucleotide that is expressed in the cell and that encodes the polypeptide.

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67. (Withdrawn-Currently amended) A method according to claim 66, further comprising: comparing (i) the quality of the corresponding phenotype immune response displayed by a test organism or part thereof mammal to which a synthetic construct comprising a tandem repeat of the first codon was provided; and (ii) the quality of the corresponding phenotype displayed by a test organism or part thereof mammal to which a synthetic construct comprising a tandem repeat of a second codon was provided, wherein the second codon encodes the same amino acid as the first codon, to thereby determine the phenotypie immune response preference of the first codon relative to the phenotypie immune response preference of the second codon in the test mammal organism or part.

68. - 69. (Canceled).

- 70. (Withdrawn-Currently amended) A method according to claim 66, further comprising: introducing the synthetic construct into a selected cell of the test organism or part thereof mammal.
 - 71. -73. (Canceled).

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74. (Withdrawn-Currently amended) A method of modulating the quality of an immune response selected phenotype that is displayed by an organism a mammal of interest or part thereof- and that results from the expression of a parent polynucleotide that encodes the polypeptide, the method comprising: introducing into the organism-or-part mammal a synthetic polynucleotide that is distinguished from the parent polynucleotide by the replacement of a first codon in the parent polynucleotide with a synonymous codon that exhibits a different phenotypie immune response preference than the first codon in the mammal organism-or-part, wherein the selected phenotype is other than a phenotype conferred upon a cell-by-a polynucleotide that is expressed in the cell and that encodes the polypeptide the synthetic polynucleotide is constructed according to the method of claim 39.

75. (Withdrawn-Currently amended) A method of enhancing the quality of an immune response selected phenotype that is displayed by an organism a mammal of interest or part thereof and that results from the expression of a parent polynucleotide that encodes the polypeptide, the method comprising: introducing into the organism or part mammal a synthetic polynucleotide that is distinguished from the parent polynucleotide by the replacement of a first codon in the parent polynucleotide with a synonymous codon that exhibits a higher phenotypie immune response preference than the first codon in the mammal organism or part, wherein the selected phenotype is other than a phenotype conferred upon a cell by a polynucleotide that is expressed in the cell and that encodes the polypeptide the synthetic polynucleotide is constructed according to the method of claim 55.

76. (Withdrawn-Currently amended) A method of reducing the quality of an immune response -selected phenotype that is displayed by an organism a mammal of interest or part thereof and that results from the expression of a parent polynucleotide that encodes the polypeptide, the method comprising: introducing into the organism or part mammal a synthetic polynucleotide that is distinguished from the parent polynucleotide by the replacement of a first codon in the parent polynucleotide with a synonymous codon that exhibits a lower-phenotypie immune response preference than the first codon in the mammal organism or part, wherein the selected phenotype is other than a phenotype conferred upon a cell by a polynucleotide that is expressed in the cell and that encodes the polypeptide the synthetic polynucleotide is constructed according to the method of claim 57.

- 77. (New) A method according to claim 39, wherein the quality is a measure of the strength, intensity or grade of the immune response.
- 78. (New) A method according to claim 39, wherein the immune response conferred by the synthetic polynucleotide is stronger than the immune response conferred by the parent polynucleotide.